## C-cp-Ag Composite Electrodes: A New Approach for Metal Air Batteries

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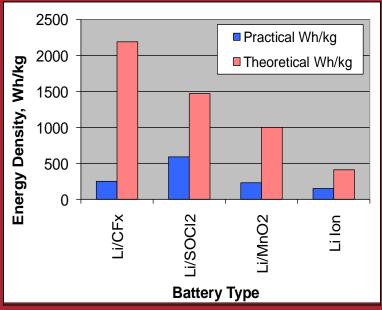
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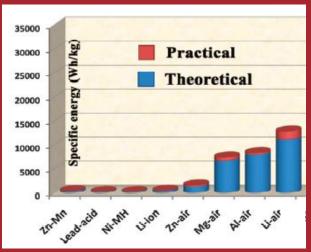
**Report Documentation Page** 

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# Metal-air batteries provide the opportunity for unprecedented energy density improvements

	theoretical			
battery	voltage (V)	capacity (Ah/kg)	energy (Wh/kg)	
Li / (CF) <sub>n</sub>	3.1	860	2,180	
Li / SOCl <sub>2</sub>	3.7	450	1,470	
Li / MnO <sub>2</sub>	3.5	310	1,010	
Li-ion	3.8	150	570	
Li / O <sub>2</sub>	3.4	3,860	13,000	
AI / O <sub>2</sub>	2.3	2,990	6,900	
Mg/O <sub>2</sub>	2.7	2,200	5,950	
Zn / O <sub>2</sub>	1.3	820	1,070	

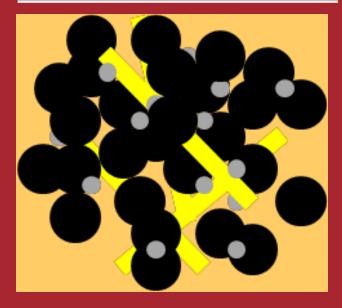




Chem. Soc. Rev., 2012, 41,2172-2192.

#### Conventional electrode fabrication

#### conventional electrode



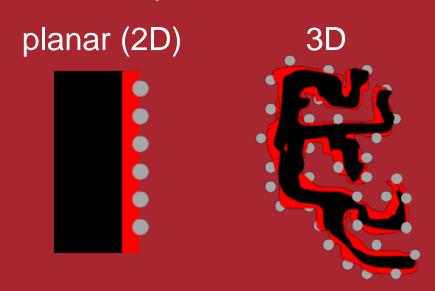
silver = active material

black = conductive additive

yellow = insulating binder

tan = metal foil current collector

#### novel composite electrode



black = conductive carbon
red = conductive polymer
silver = catalyst

## Our composite electrode strategy offers several advantages

Electrochemical deposition of layers ensures good electrical contact among composite electrode components and electrolyte.

This is a transferrable concept that can be extended to prepare 2D and 3D layered composite electrodes.

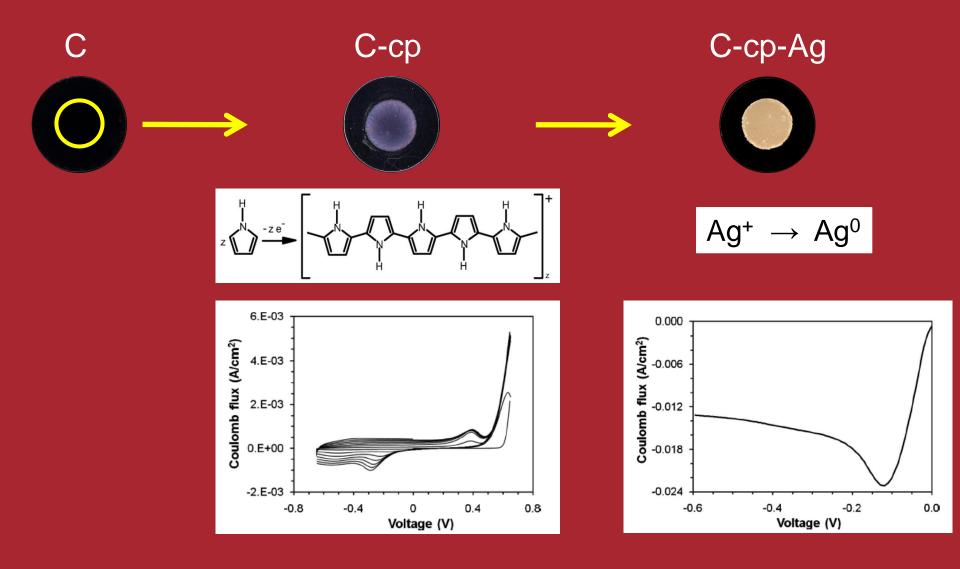
Depending on the nature of the current collector (cc), and conducting polymer (cp), morphology and porosity of the layered composite can be tuned.

Due to the conductive nature of the composite, even small quantities of catalyst (Ag) should exhibit high oxygen reduction activity.

### Project objectives

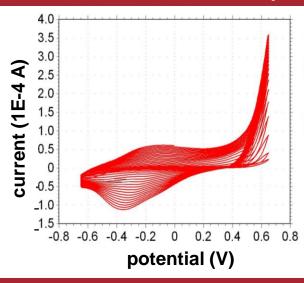
- i. Demonstrate new composite electrode based on carbon-conductive polymer-silver (C-cp-Ag) composite.
- ii. Evaluate composite oxygen reduction activity.
- iii. Assess roles of composite components.
- iv. Investigate non-aqueous oxygen reduction mechanism.
- v. Develop and investigate 3D C-cp-Ag composite.

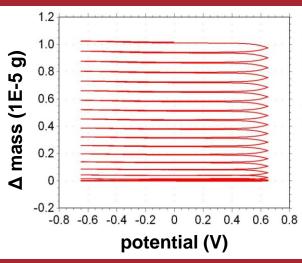
## Electrodeposition is desirable for fabrication

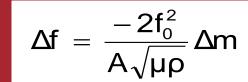


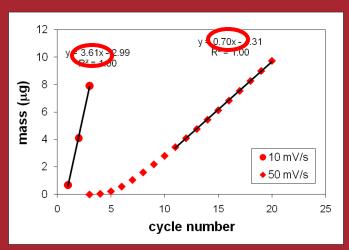
### C-cp-Ag prep can be quantitatively controlled

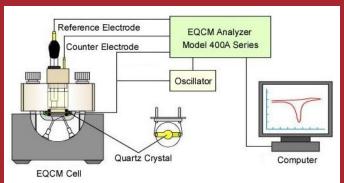
#### cp deposition





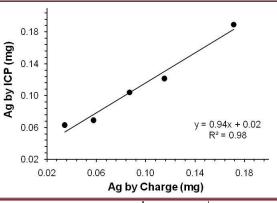




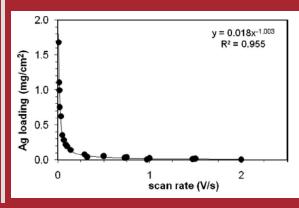


#### Ag deposition

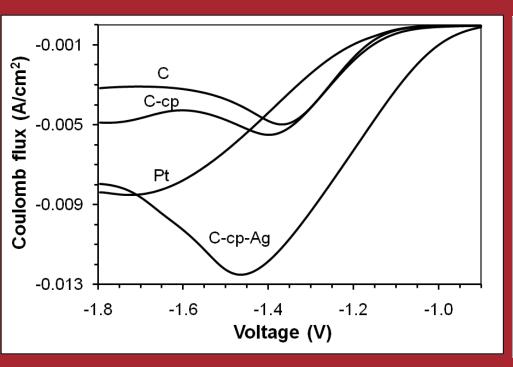
$$q = n \cdot F$$

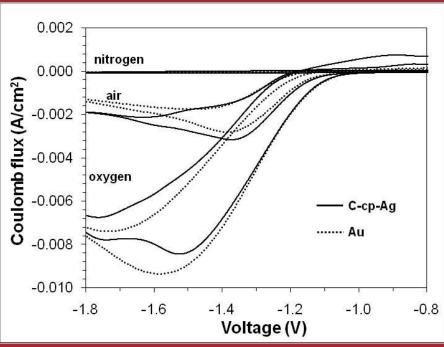


	trial 1	trial 2
EQCM mass (μg)	19.0	5.94
Faraday mass (μg)	21.7	6.75
Δ (%)	12.5	12.0



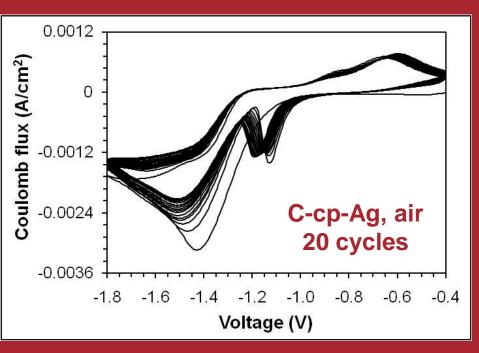
## C-cp-Ag compares favorably to benchmarks

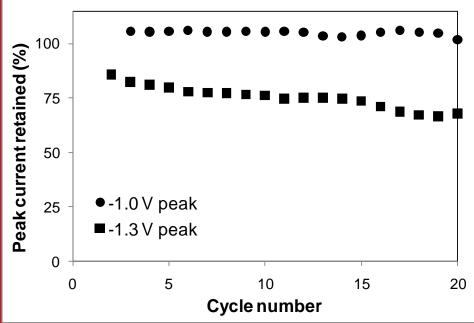




C-cp-Ag composite shows activity
2.6x higher than C
1.4x higher than Pt
comparable to Au

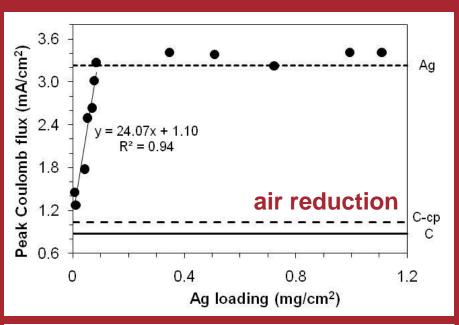
### C-cp-Ag composite shows catalytic activity

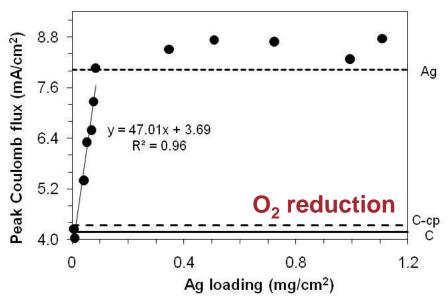




C-cp-Ag composite electrodes retain high oxygen reduction capability on multiple cycling.

## Low Ag loading is required for optimization





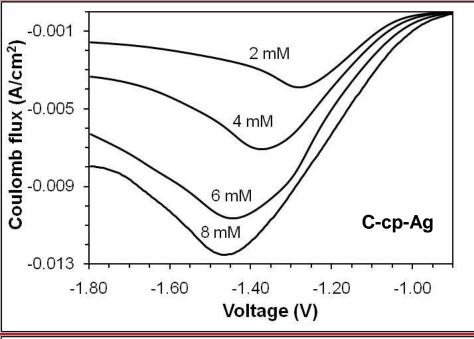
response optimized at Ag loading of 0.08 mg/cm<sup>2</sup>

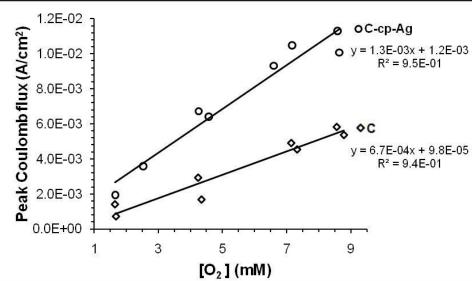
activity in air was 20 – 40% of the activity in pure oxygen

using density of Ag metal, suggests that 80 nm is minimum Ag thickness required

#### kinetics:

#### C-cp-Ag shows well behaved ORR



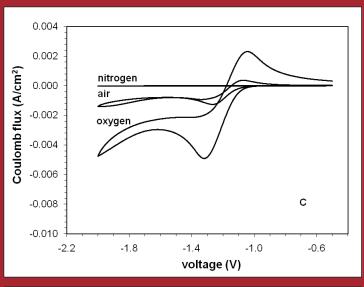


linear response to [O<sub>2</sub>], consistent with 1<sup>st</sup> order process

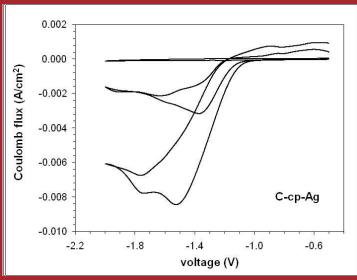
C-cp-Ag shows enhanced activity at all [O<sub>2</sub>]

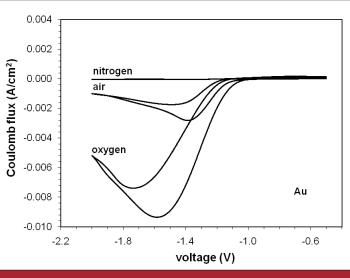
Ag loading of 0.3 mg/cm<sup>2</sup> for C-cp-Ag composite

## C-cp-Ag and Au-cp-Ag show similar profiles

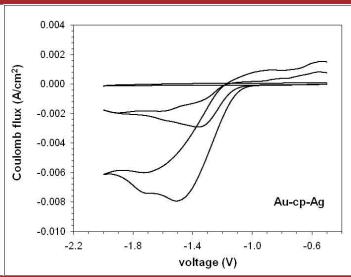


cp, then Ag deposition

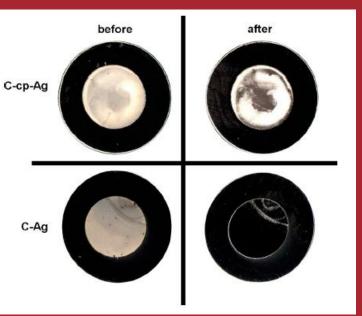


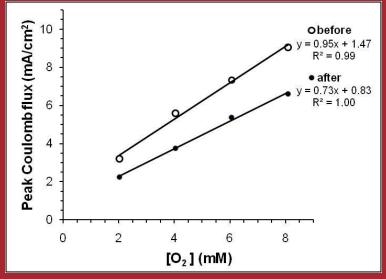


cp coating mitigates impact of substrate



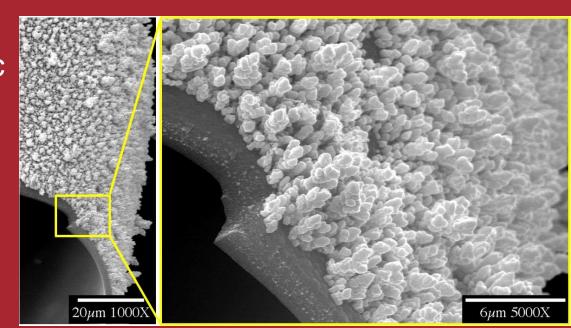
## C-cp-Ag composites are robust



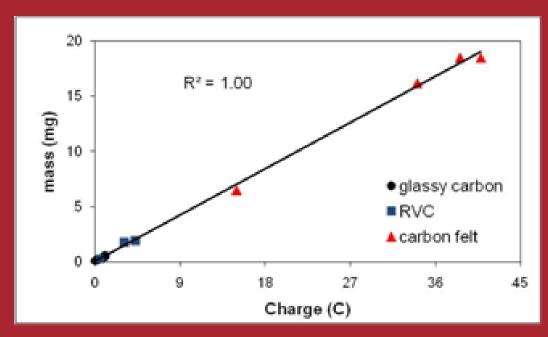


number	12		
average	73%		
median	72%		
st. dev.	8.7%		

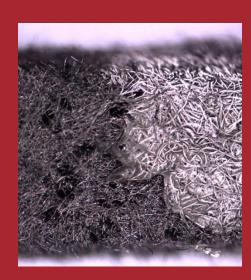
demonstrated macroscopic and microscopic Ag (and activity) retention after physical stresses



### 3D C-cp-Ag electrodes can be fabricated



C-felt

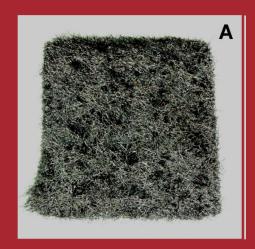


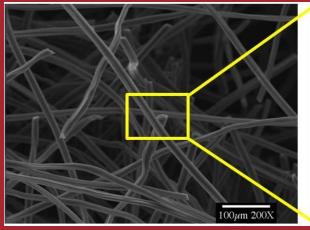
Achieved successful cp and Ag-cp deposition on non-planar (3D) substrates of varying geometry and porosity.

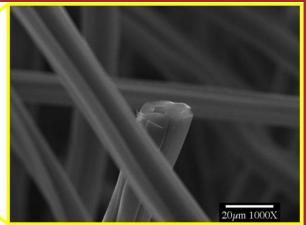
RVC



### 3D C-cp-Ag electrodes can be fabricated







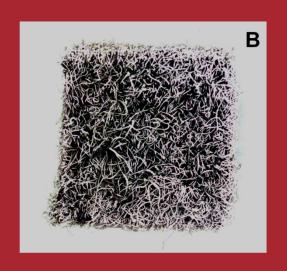
electrode	electrode surface area (cm²)			
electi ode	reported	planar	geometric	electrochemical
С	0.071	0.069	0.069	0.067
Cfelt	121	1.0	3.2	17

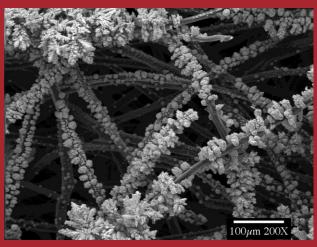
estimated cp thickness ~4X greater for C than for Cfelt

electrode	estimated cp	electrode surface area (cm²)			
electi ode	thickness (µm)	planar	geometric	electrochemical	
C-cp	3.5	0.076	0.076	0.053	
Cfelt-cp	0.9	1.0	3.2	9.6	

electrochemically active surface area reduced 20% for C, > 40% for Cfelt

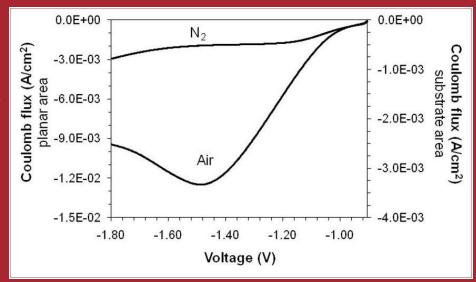
## 3D C-cp-Ag composite electrodes



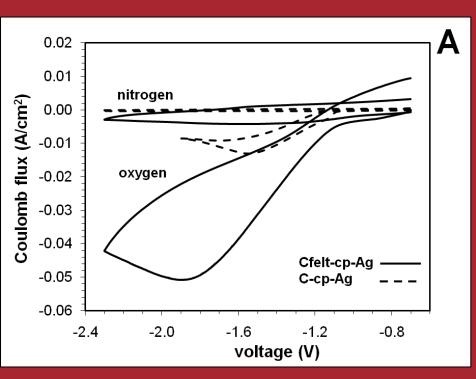


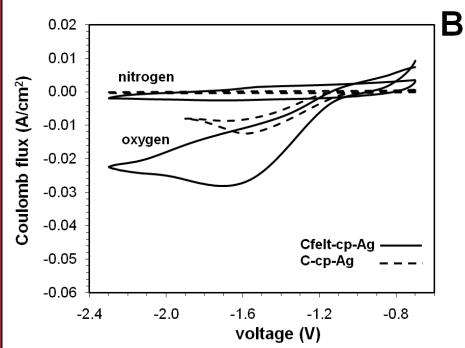
For C-cp-Ag composites, use of 3D substrates provided 4X improvement over 2D substrates based on planar area.

There is opportunity to further improve homogeneity of the 3D composite.



## 3D C-cp-Ag composite electrodes





Cycle 1 – 2 peak Coulomb flux was unchanged for the planar C-cp-Ag electrode, while the 3D Cfelt-cp-Ag composite showed a significant decrease.

Difference is consistent with reduction on the surface and interior during cycle 1, with surface only on cycle 2.

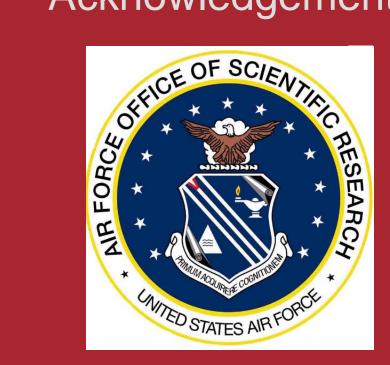
#### **Publications**

- 1) "Metal-air electrochemical cells: Silver-polymer-carbon composite air electrodes." *Electrochem. S. S. Lett.* **2010,** 13(11), A162-A164.
- 2) "Electrodes for nonaqueous oxygen reduction based upon conductive polymer-silver composites." *J. Electrochem. Society.* **2011,** *158(3),* A223-A226.
- 3) "Oxygen reduction activity of carbon-conductive polymer-silver composite electrodes." ECS Transactions, **2012**, *41*(10), 9-13.
- 4) "Three dimensional carbon-conductive polymer-silver (C-cp Ag) composite electrodes for metal-air batteries." *J. Composite Materials. in press.*
- 5) "Secondary Battery Science: At the Confluence of Electrochemistry and Materials Engineering." *Electrochemistry*. *in press*. (invited highlight)
- 6) "Mechanistic investigation of the oxygen reduction reaction on carbon-conductive polymer-silver composites." *in preparation.*
- 7) "Activation energy of oxygen reduction on carbon substrates in nonaqueous media." *in preparation.*

#### **Patent**

"Electrodes for metal-air batteries and fuel cells." US 2010-300719P, Feb 2, 2010; WO 2011097286, Aug. 11, 2011; WO 2011097286, Dec. 29, 2011.

### Acknowledgement



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